

**REFUSAL TO CONSENT to REPEAT NEWBORN SCREENING**I/We, \_\_\_\_\_, the parent(s) of  
Name of parent(s)\_\_\_\_\_, born on \_\_\_\_\_ at  
Infant's name Date of birth\_\_\_\_\_, refuse to have blood taken from our child for the purpose of  
Place of birth

determining if (s)he might have a metabolic or other disorder. We understand that the initial specimen obtained was unsatisfactory for testing or indicated a need to repeat the screening. The conditions tested for include the 29 conditions listed below. I/we understand that the Vermont Department of Health recommends all babies be tested for these conditions in the newborn period.

<i>3-Methylcrotonyl-CoA carboxylase deficiency (3MCC)</i>	<i>Maple syrup urine disease (MSUD)</i>
<i>3-OH 3-CH<sub>3</sub> glutaric aciduria (HMG)</i>	<i>Medium-chain acyl-CoA dehydrogenase deficiency (MCAD)</i>
<i>Argininosuccinic acidemia (ASA)</i>	<i>Methylmalonic acidemia (Cbl A, B)</i>
<i>Beta-ketothiolase deficiency (BKT)</i>	<i>Methylmalonic acidemia (mutase deficiency) (MUT)</i>
<i>Biotinidase deficiency (BIOT)</i>	<i>Multiple carboxylase deficiency (MCD)</i>
<i>Carnitine uptake defect (CUD)</i>	<i>Phenylketonuria (PKU)</i>
<i>Citrullinemia (CIT)</i>	<i>Propionic acidemia (PROP)</i>
<i>Congenital adrenal hyperplasia (CAH)</i>	<i>Severe Combined Immunodeficiency (SCID)</i>
<i>Congenital hypothyroidism (HYPOTH)</i>	<i>Sickle cell anemia (SCA)</i>
<i>Cystic fibrosis (CF)</i>	<i>Trifunctional protein deficiency (TFP)</i>
<i>Galactosemia (GALT)</i>	<i>Tyrosinemia type I (TYR I)</i>
<i>Glutaric acidemia type I (GA I)</i>	<i>Very long-chain acyl-CoA dehydrogenase deficiency (VLCAD)</i>
<i>Hb S/Beta-thalassemia (Hb S/Th)</i>	
<i>Hb S/C disease (Hb S/C)</i>	
<i>Homocystinuria (HCY)</i>	
<i>Isovaleric acidemia (IVA)</i>	
<i>Long-chain L-3-OH acyl-CoA dehydrogenase deficiency (LCHAD)</i>	

*Other types of screening tests which can be done in the home or hospital include screening for hearing loss and pulse oximetry to screen for Critical Congenital Heart Disease.*

- I/we have read the brochure provided by the Vermont Department of Health (VDH) Newborn Screening Program and understand that VDH recommends that all babies be tested for these conditions in the newborn period.
- I/we understand that the VT Newborn Screening Program recommends that follow-up testing be done as a result of the specimen which was obtained \_\_\_\_\_ and revealed \_\_\_\_\_.
- I/we feel that we have all the information necessary and have made the decision **not** to have newborn screening repeated for our baby/babies.
- I/we do not wish to discuss newborn screening further with newborn screening staff, our baby's doctor, or other care providers who are available to answer related questions.
- I/we understand that if our baby does have one of these conditions and it is not diagnosed in the newborn period, the risk that our child could have health problems, including intellectual disabilities and/or death, could be very high.

\_\_\_\_\_  
Signature of parent(s) date\_\_\_\_\_  
Signature of witness date

Please mail this form to the Vermont Newborn Screening Program, PO Box 70, 108 Cherry St., Burlington, VT 05402. Call (802) 951-5180 with questions. Rev. 11/03/16